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Attorney Docket No. 6413.204-US Ebdrup et al. Serial No. 10/614,233 Filed July 7, 2003

**CLAIM LISTING** 

What is claimed is:

1. (Currently amended) A method of:

- a) inhibiting the lipolytic activity of hormone-sensitive lipase against triacylglycerols, diacylglycerols, cholesterol acyl esters or steroid acyl esters; and/or
- b) modulating the plasma level of free fatty acids, glycerol, LDL-cholesterol, HDLcholesterol, insulin and/or glucose; and/or
- c) modulating intracellular triacylglycerol and cholesterol ester stores, intracellular level of fatty acids, fatty acid esters, diacylglycerols, phosphatidic acids, long chain acyl-CoA's as well as citrate or malonyl-CoA; and/or
- d) increasing insulin sensitivity in adipose tissue, skeletal muscle, liver or pancreatic β cells;
- e) modulating insulin secretion from pancreatic β cells; and/or
- f) inhibiting male fertility

in a patient comprising, administering to a patient in need of such method a therapeutically effective amount of a boronic acid, an ester thereof, a prodrug thereof,

wherein the boronic acid, an ester thereof or a prodrug thereof is of the general formula I

$$R^3 \longrightarrow B \qquad (I)$$

$$O \longrightarrow R^2$$

wherein R<sup>1</sup> and R<sup>2</sup> are independently selected from hydrogen, C<sub>1-6</sub>-alkyl, C<sub>2-6</sub>-alkenyl, alkynyl, aryl, heteroaryl, C3-8-heterocyclyl and C3-10-cycloalkyl, wherein each of C1-6-alkyl, C2. 6-alkenyl, C2-6-alkynyl, aryl, heteroaryl, C3-8-heterocyclyl and C3-10-cycloalkyl is optionally substituted with one or more substituents independently selected from hydroxy, sulfanyl, sulfonyl, sulfinyl, oxo, thioxo, halogen, amino, imino, cyano, nitro, silyl, boranyl, phosphinyl, selenyl, germyl, C1-6-alkyl, C2-6-alkenyl, C2-6-alkynyl, aryl, heteroaryl, C3-8heterocyclyl and C3-10-cycloalkyl, wherein each of hydroxy, sulfanyl, sulfonyl, sulfinyl, amino, imino, silyl, boranyl, phosphinyl, selenyl, germyl, C1-6-alkyl, C2-6-alkenyl, C2-6-